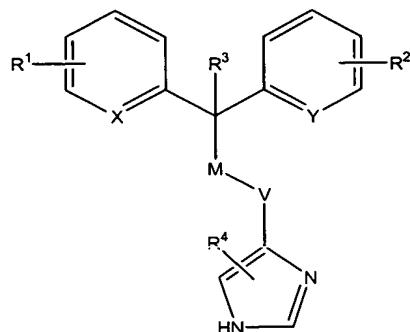


The listing of claims will replace all prior versions and listing of claims in the application:

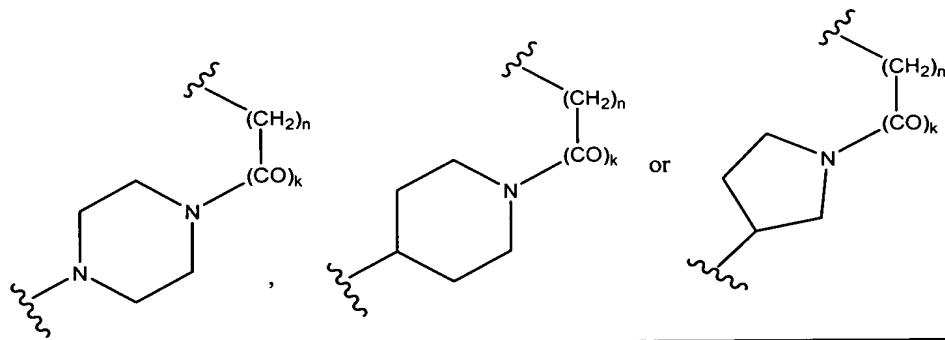
Listing of Claims:

Claim 1'(currently amended): A compound, or enantiomers, stereoisomers and tautomers thereof, or pharmaceutically acceptable salts or solvates of said compound, with said compound having the general structure shown in Formula I:

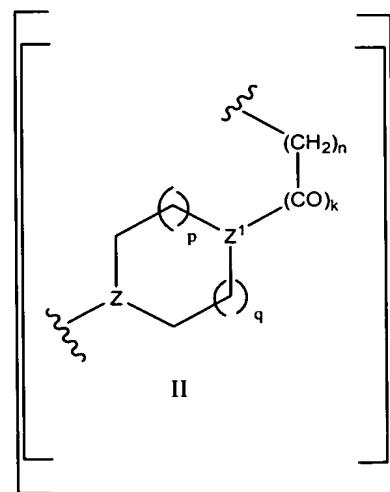


Formula I

wherein M is a moiety of the formula:



having a general structure shown in Formula II:



where $k = 0$ or 1 , and $n = 0$ - 5 [[, and $p = q = 0$, 1 or 2]];

V is a moiety selected from the group consisting of C_1 - C_8 alkyl;

$-(CH_2)_x-A-(CH_2)_y-$; and $-(CH_2)_c-A-(CH_2)_m-C(O)-N(R^7)-(CH_2)_d-$, where A is $-O-$, $-S(O)r-$, and $-NR^7-$;

$m = 0$, 1 , 2 or 3 ; x is a whole number in the range 2 - 8 ; y is a whole number in the range 1 - 5 ; c is a whole number in the range 2 - 4 ; and $r = 0$, 1 or 2 ; d is a number in the range 0 - 5 ;

[[X and Y are independently selected from the group consisting of N , and CH ;]]

one of X is N and the other is CH :

[[Z and Z' can be the same or different, each being independently selected from the group consisting of N , CH and $N(O)$;]]

R^1 and R^2 may each number 1 - 4 and are independently selected from the group consisting of hydrogen, lower alkyl, lower alkoxy, halogen, polyhalolower alkyl, polyhalolower alkoxy, $-OH$, CN , NO_2 , or $COOR^8$;

R^3 is selected from hydrogen, lower alkyl, lower alkoxy, hydroxyl, ~~with the proviso that when n and k are both 0, then R^3 is not $-OH$ or alkoxy;~~

R^4 is selected from the group consisting of hydrogen, lower alkyl, polyhalolower alkyl or $-OH$; and

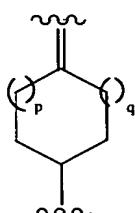
R^7 and R^8 are independently selected from hydrogen, lower alkyl, substituted or unsubstituted phenyl; and substituted or unsubstituted benzyl, wherein said term "substituted" means optional substitution from one or more moieties selected from the group consisting of alkyl, alkoxy, $-CF_3$, halogen or aryl.

Claim 2 (original): The compound of claim 1, wherein R^4 is H .

Claim 3 (original): The compound of claim 2, wherein R^1 and R^2 are independently selected from H , halogen, or polyhalolower alkyl.

Claim 4 (canceled).

Claim 5 (amended): The compound of claim 1, wherein M is a piperazine. [[:



and p = q = 1.]]

M is piperidine,
C Claim 7¹⁸ (original): The compound of claim 4, wherein R⁴ is H; R¹ = R² = H, halogen, hydroxy or alkoxy; and R³ is H or lower alkyl.

B' Claim 8¹⁸ (original): The compound of Claim 6, wherein V = C₁ – C₈ alkyl.

Claim 9¹⁸ (original): The compound of claim 5, wherein R⁴ is H; and R¹ = R² = H, halogen, hydroxy or alkoxy.

Claim 10¹⁸ (original): The compound of Claim 8, wherein V is C₁ – C₈ alkyl.

Claim 11¹⁸ (previously amended): A pharmaceutical composition comprising as an active ingredient a compound of claim 1 and a pharmaceutically acceptable carrier.

Claim 11 (previously canceled).

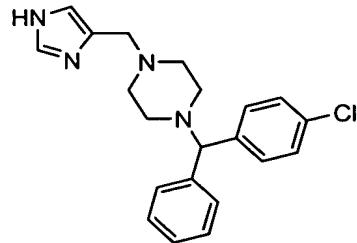
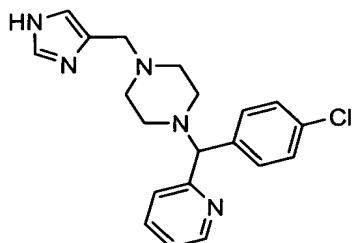
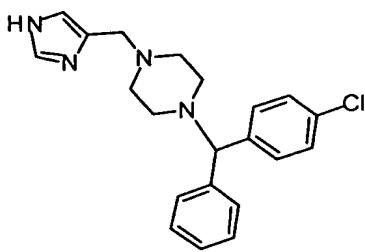
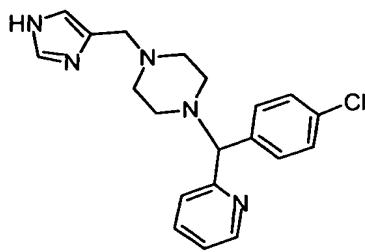
Claim 12 (previously canceled).

Claim 13¹⁸ (currently amended): A method of treating airway and gastrointestinal disorders ~~inflammation, allergy, nasal congestion, diseases of the GI tract, cardiovascular disease, or disturbances of the central nervous system as well as allergy-induced airway responses, and obesity~~, said method comprising administering to a mammalian patient in need of such treatment a pharmaceutical composition which comprises therapeutically effective amounts of a compound of claim 1.

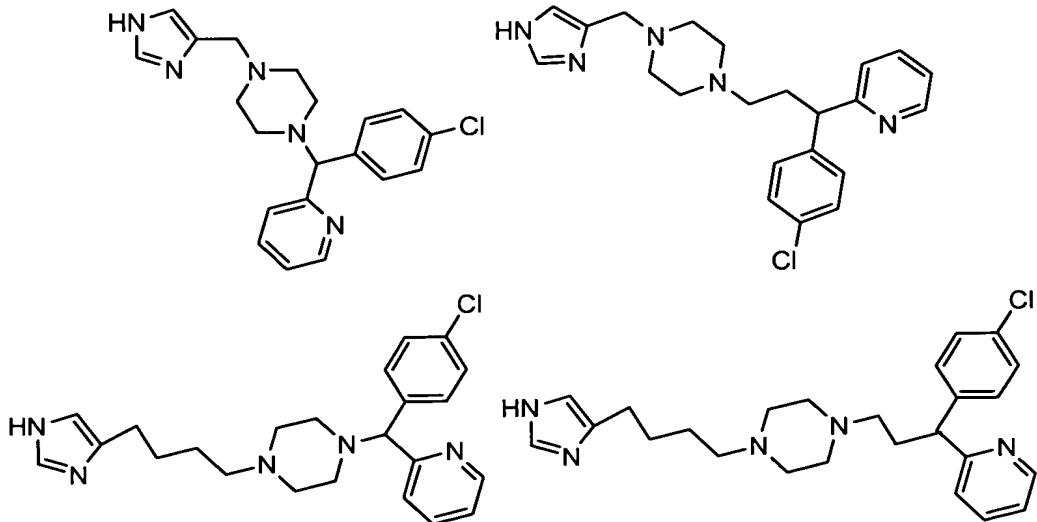
Claim 14 (previously canceled).

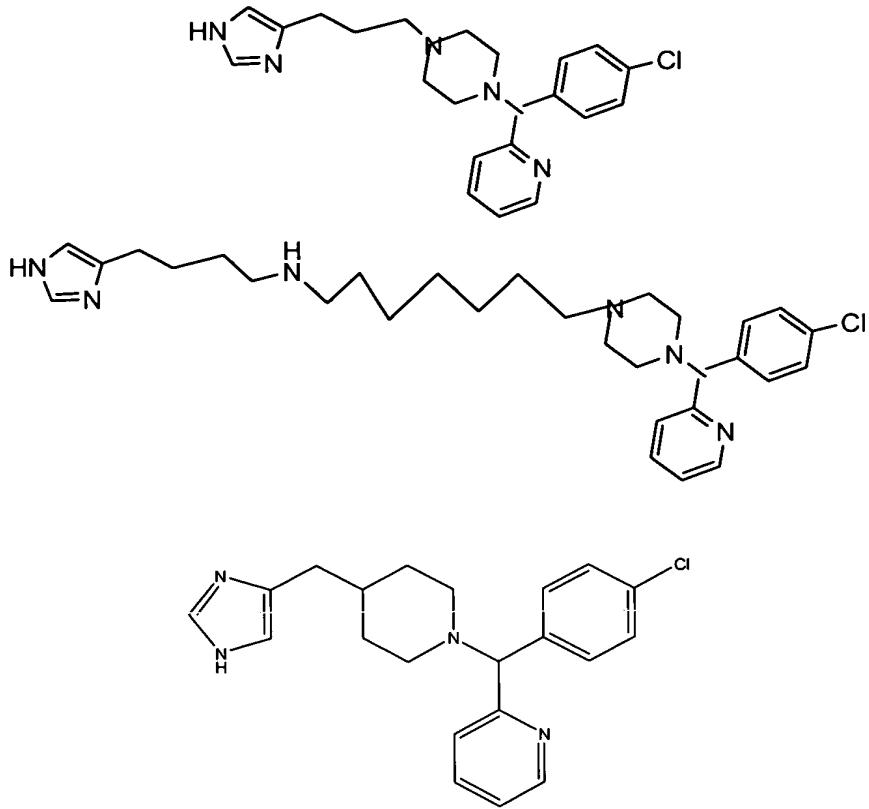
Claim 15¹⁸ (currently Amended): A method of preparing a pharmaceutical composition for treating airway and gastrointestinal disorders ~~inflammation, allergy, nasal congestion, diseases of the GI tract, cardiovascular disease, or disturbances of the central nervous system as well as allergy-induced airway responses, and obesity~~, said method comprising bringing into intimate contact a compound of claim 1 and a pharmaceutically acceptable carrier.

Claim 16¹⁸ (currently amended): A compound exhibiting H₃ antagonist activity, or enantiomers, stereoisomers and tautomers of said compound, or pharmaceutically acceptable salts or solvates of said compound, said compound being selected from the compounds with structures listed below:



Claim 17 (currently amended): A compound exhibiting both H₁ and H₃ antagonist activity, or enantiomers, stereoisomers and tautomers of said compound, or pharmaceutically acceptable salts or solvates of said compound, said compound being selected from the compounds with structures listed below:





Claim 18 (currently amended): A pharmaceutical composition for treating airway and gastrointestinal disorders ~~inflammation, allergy, nasal congestion, diseases of the GI tract, cardiovascular disease, or disturbances of the central nervous system as well as allergy-induced airway responses, and obesity,~~ said composition comprising therapeutically effective amount of a compound of claim 16 or claim 17 and a pharmaceutically acceptable carrier.

Claims 19-21 (canceled).

Claim 22 (new): The compound of claim 1, wherein M is a piperidine.

Claim 23 (new): The compound of claim 1, wherein M is a pyrrolidine.